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## Amendments to the Specification:

This listing of claims will replace all prior versions and listings of claims in the application:

1. (Previously presented) A method of generating a protein array, the method comprising:

(a) inserting a marker DNA sequence in frame immediately following a start codon of

each of a plurality of target DNA sequences or immediately preceding a stop codon of each of a

plurality of target DNA sequences or both, to form a plurality of modified DNA sequences which

encode a plurality of modified amino acid sequences each comprising a marker moiety;

(b) expressing the plurality of modified amino acid sequences from the plurality of

modified DNA sequences;

(c) bringing the plurality of modified amino acid sequences into contact with a solid

support wherein the marker moiety of the plurality of modified amino acid sequences is able to

attach to the solid support, thereby generating a protein array, and

(d) washing said solid support to remove unbound proteins.

2. (Previously presented) The method as claimed in claim 1 wherein the marker moiety is a

peptide sequence selected from the group consisting of:

(a) a histidine tag;

(b) a complete protein or protein domain; and

(c) a maltose binding protein domain.

3. (Previously presented) The method as claimed in claim 1 wherein the marker moiety

allows for purification of the individual proteins in the array.

4. (Previously presented) The method of claim 1 wherein the marker DNA sequence is

inserted such that the start or stop codon for each of the proteins is replaced.

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- 5-7. (Canceled)
- 8-12. (Withdrawn)
- 13. (Previously presented) A method of generating an antibody array which comprises
- (a) bringing a protein array, made according to any one of claims 1 to 4, into contact with an antibody library, such that one or more proteins in the protein array bind to at least one antibody in the antibody library;
  - (b) removing any unbound antibodies; and
  - (c) immobilisation of those antibodies bound to proteins in the protein array.
- 14. (Withdrawn)
- 15. (Canceled)
- 16. (Previously presented) The method of claim 1 wherein the marker DNA sequence is inserted immediately preceding a stop codon of a target DNA sequence by:
- (a) digesting the target DNA sequence such that it has a 5' overhang wherein the stop codon is comprised in the first three nucleotides counting from the 3' side of the overhang;
- (b) annealing the marker DNA sequence to the overhang wherein the marker DNA sequence comprises a sequence complementary to the first four nucleotides of the overhang counting from the 3' side;
  - (c) ligating the marker DNA sequence to the target DNA sequence.
- 17. (Previously presented) The method of claim 1 wherein the marker DNA sequence is inserted immediately following a start codon of a target DNA sequence by;
- (a) digesting the target DNA sequence such that it has a 5' overhang wherein the start codon is comprised in the first three nucleotides counting from the 3' side of the overhang;

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(b) annealing the marker DNA sequence to the overhang wherein the marker DNA

sequence comprises a sequence complementary to the first four nucleotides of the overhang

counting from the 3' side;

(c) ligating the marker DNA sequence to the target DNA sequence.

18. (Previously presented) The method of any one of claims 1 to 4 wherein the protein array

comprises serine proteases, kinases or p450 enzymes.

19. (Previously presented) The method of any one of claims 1 to 4 wherein said plurality of

modified amino acid sequences are modified human amino acid sequences.

20. (Previously presented) The method of claim 1 wherein the marker moiety is selected

from the group consisting of FLAG and Strep.

21. (Previously presented) The method of claim 1 or 2 wherein the marker moiety is post-

translationally modified.

22. (Previously presented) The method of claim 21 wherein the post-translational

modification comprises the addition of a biotin or a lipid molecule.

23. (Previously presented) The method of claim 1 wherein said modified amino acid

sequences are folded into the correct conformation.

24. (Previously presented) The method of claim 1 wherein said inserting step inserts a

marker DNA sequence in frame immediately following a start codon of each plurality target

DNA sequence and immediately preceding a stop codon of each of a plurality of target DNA

sequences, to form a plurality of modified DNA sequences which encode a plurality of modified

amino acid sequences each comprising two marker moieties.

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25. (Withdrawn)

26. (Currently amended) A method of screening for antibodies which recognize each protein in the array, the method comprising:

- (a) contacting the antibodies with a spatially defined array comprising a plurality of array bound proteins produced according to any one of claims 1-4, with each array bound protein being at a different position on a solid support, wherein the plurality of array bound proteins comprises a plurality of different proteins expressed in a single species; and
  - (b) detecting any interaction between the array bound proteins and the antibodies.
- 27. (New) The method of claim 1, wherein the marker moiety provides a high-affinity attachment to the solid support.